

JAKLEWICZ, Hanna

Psychiatric and social conditioning in juvenile crimes.
Neurol.neurochir. psychiat. Pol. 14 no. 2:303-307 Mr-Apr '64.

1. Z Kliniki Chorob Psychiczych AM w Gdansk (Kierownik:
prof. dr T.Bilikiewicz).

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JAKLEWICZ, Przemyslaw, mgr inz.; KUPRAS, Krystyn, mgr inz.

Designing ship's ordinate lines by means of electronic computers. Bud
okretowe Warszawa 8 no.3:81-85 Mr '63.

1. Centralne Biuro Konstrukcji Okretowych Nr 1, Gdansk.

JAKLIC, OTMAR

YUGOSLAVIA/Engineering - Electric Power Apr/May 49
Stations
Construction

"Hydromechanical Equipment of Pillar-Type Electric
Power Stations," Otmar Jaklic, Engr Maribor 4 1/2 pp

"Elektrotehnicki vesnik" No 4/5

Pillar-type power-station construction is becoming
more common, and hydromechanical equipment must be
made to conform with it. Describes equipment of
"Mariborski otok" station in some detail, with
examples of cooperation received from various
enterprises in manufacture of large machine elements.
Includes twelve illustrations.

150R26

JAKLINSKI, Adam

Natural death or death by injury of shock susceptible body parts.
Arch.med.sad., Warszawa 6:79-81 1955.

1. Z Zakladu Medycyny Sadowej A.M. w Lublinie. Kierownik: prof.
dr. W. Dzulynski.

(WOUNDS AND INJURIES

heart region after accid. fract. of ribs & sternum,
causing sudden death, medicolegal determ. by post-
mortem exam.)

(DEATH, SUDDEN,

caused by trauma of heart region after accid. fract.
of ribs & sternum, medicolegal determ. by post-mortem
exam.)

(ACCIDENTS

fract. of ribs & sternum causing inj. of heart region
& sudden death, medicolegal determ. of cause of death)

JAKLINSKI, Adam.

An unusual case of traumatic gangrene of the lungs. Arch.med.
sad., Warszawa 6:82-86 1955.

1. Z Zakladu Medycyny Sadowej A.M. w Lublinie. Kierownik:
prof. dr W. Dzulynski.

(LUNGS, gangrene

caused by inj., fatal, post-mortem determ. of cause
of death, medicolegal aspect)

(WOUNDS AND INJURIES,

lungs causing gangrene & death, post-mortem determ.
medicolegal aspects)

(GANGRENE,

lungs, caused by inj, fatal, post-mortem determ. of
cause of death, medicolegal aspects.)

GERKOWICZ, T.; JAKLINSKI, A.

Case of endomyocardial fibroelastosis. *Pediat. polska* 31 no.4:
445-448 Apr 56.

1. Z Kliniki Chorob Dzieci A.M. w Lublinie. Kier.: doc. dr. med.
W. Klepacki i z Zakladu Medycyny Sadowej A.M. w Lublinie Kier.:
prof. dr. med. W. Dzulyński, Lublin, Staszica 11, Klin. Ped.
(CARDIAC ENLARGEMENT, in infant and child,
endocardial fibroelastosis (Pol))

JAKLINSKI, Andrzej, doc. dr.; ERYC, Stanislaw

Evaluation of sequelae of injury in deformative fibrous bone degeneration (Paget's osteitis deformans). Pol. tyg. lek. 20 no.3:108-110 18 Ja '65

1. Z Zakladu Medycyny Sadowej Akademii Medycznej w Lublinie (Kierownik: doc. dr. A. Jaklin-'''' - Zakladu Radiologii Akademii Medycznej w Lublinie (Kierownik: doc. dr. K. Skorzynski).

POLAND

JABLINSKI, Andrzej, Department of Legal Medicine (Zaklad Medycyny Sadowej), AM [Akademia Medyczna, Medical Academy] in Lublin (Director: Prof. Dr. W. DZULYNSKI)

"Experimental Studies on Cerebrospinal Fluid Chlorides Concentration in Post-Mortem Examinations."

Warsaw, Polski Tygodnik Lekarski, Vol 17, No 39, 24 Sep 62, pp 1499-1502.

Abstract: [Author's English summary modified] CSF from terminal and large reservoir were studied by Mohr method for chloride ion concentration 2-96 hours after death on 52 bodies. Correlation coefficient of $r=0.26$ established for large, and none for terminal reservoir CSF. Test cannot be used to establish time of death. Of 10 references, 6 are in the English, 4 in the German, and 2 in the Polish language.

1/1

JAKLOVA, Stanislava, inz.

Blast furnace operation control by measurement of pressure differences. Hut listy 19 no. 4: 268-271 Ap '64.

1. Research and Testing Institute, Nova hut Klémenta Gottwalda, Ostrava-Kuncice.

JAKLOVESKY, A.

Results obtained with a new anti-diarrhoeic dietetic product,
cellulose-lignin powder. Rumanian M. Rev. 3 no.4:30-32 O-D '59.

1. Department of Paediatrics of the Unified District Hospital in
Oradea.

(DIARRHEA, in inf. & childh.)

(LIGNIN, therapy)

(CELLULOSE, therapy)

2.110A 2.
Given Data

Country: Romania

Academic Degrees: Dr.

Affiliation: #)

Source: Bucharest, Microbiologia, Parasitologia, Epidemiologia, No 3,
May-Jun 61, pp 251-262.

Date: "Data Concerning the Appearance of Resistance to Chloramphenicol of Some Sh. Flexneri Strains and the Testing of Their Immuno-genicity."

Co-authors:

HADMACY, C. Dr.

JAKLOVICH, A., DT.

Work performed at Clinic No 2 of Tg. Mures/and at the Laboratory of the Odorhei National Sanepid (Laboratorul Sanepidului National Odorhei).

JAKLOVSZKY, Alfons

Notes on catamnesis of cases of Bouillaud-Sokolski's rheumatism hospitalized
in the children's clinic of Odorhei between 1950-1955. Probl. reumat.,
Bucur. no.5:127-129 1958.

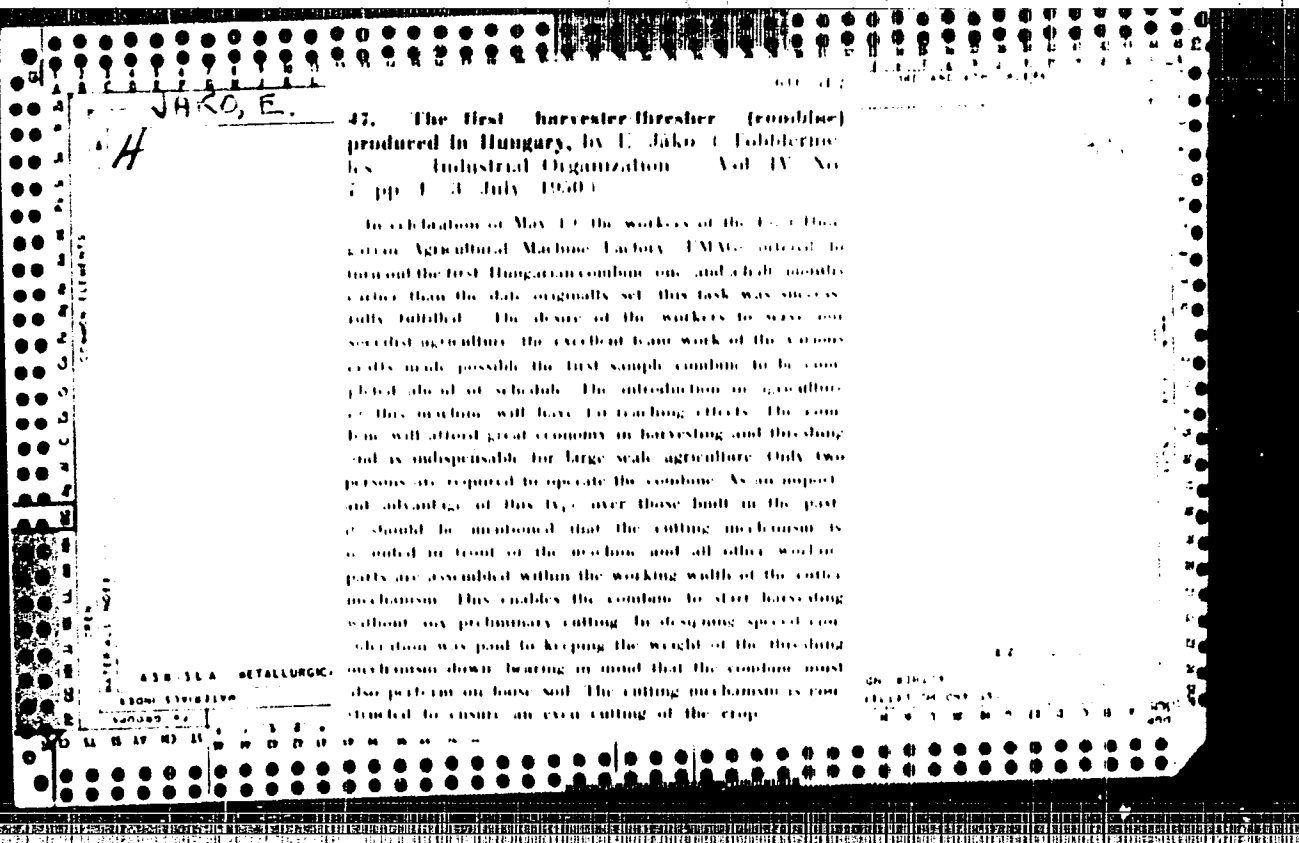
(RHEUMATIC HEART DISEASE

evolution & results of ther. in child. of Odorhei, Rumania)

JAKLOWICZ, DAZIMIERZ.

Obsługa radiotelefonu i echosondy. Warszawa, wydawn. komunikacyjne,
1954. 81 p. (Poradnik rybaka morskiego, zesz. 8)

SOURCE: East European Accession List (EEAL) Library of Congress
Vol. 5, no. 8, August 1956



JAKO, F.

MEZOGAZDASGI IPAR — AGRICULTURAL INDUSTRY
Vol. IV — 1950
No. 9, Sept.

31

F. Jakó

617111

Budding and equipping stumps

pp. 11-19

ASH 51.1 METALLURGICAL LITERATURE CLASSIFICATION

"Tej es tejtermek, hal, hasznos elvöd es lottvöd, hutes es jeggyartas.
00szellitottak: Jako Frigyes et al.) Kereskedelmi Szekkonv- es Larkiedo.
128 o. (Kereskedelmi aruismemet) (Milk and dairy products, fish, useful
game and game products, refrigeration, and the production of ice; a hand-
book on properties and methods).

SQ: East European Accessions List, Vol 3, No 8, Aug 1954.

MISSURA, Tibor, dr.; JAKO, Geza

Besnier-Boeck-Schaumann sarcoidosis of the upper respiratory tract. Orv. hetil. 96 no.20:556-557 15 May 55.

1. A Peterfy Sandor-utcai korhaz-Rendelo (igazgato-
Lendvai, Jozsef dr.) Ful- orr- gegesztalyanak (foorvon:
Fleischmann, Lasso, az Orvostudomanyok Doktora) kozlensege.
(SARCOIDOSIS,
nose.)
(NASAL CAVITY, diseases,
sarcoidosis,)

SZMUK, Imre, dr.; BACH, Imre, dr.; DANZIGER, Laszlo, dr.; FEKETE, Balazs, dr.;
FLEISCHMANN, Laszlo, dr.; JAKO, Gábor, dr.; MISSURA, Tibor, dr.;
POPPER, Szuzsanna, dr.; SZABADOS, Daisy, dr.

Use of radioiodine in localization of inflamed regions (foci,
abscesses). Orv. hetil. 97 no.34:949-951 19 Aug 56.

1. A Fovarosí Peterfy Sándor u. Korhazrendelő (igazgató:
Landvai, József, dr.) közleménye.

(BRAIN, abscess

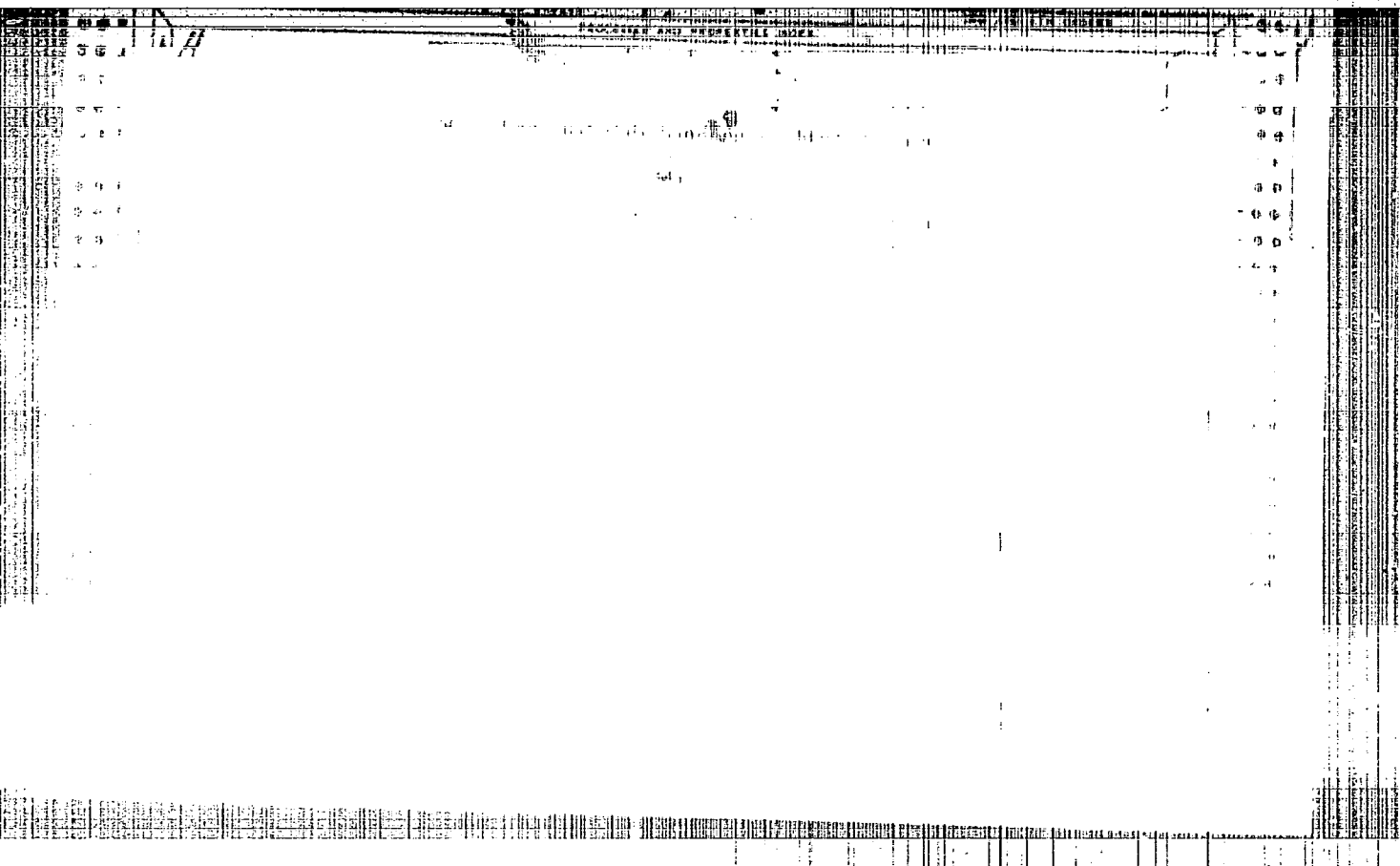
exper., localization with radioiodine in dogs (Hun))

(IODINE, radioactive

in localization of exper. brain abscesses in dogs (Hun))

"APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619420007-2



APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619420007-2"

BANYASZ, T.; JAKO, J.; HORVATH, I.

On the effect of treatment with butylbiguanide on the liver function. Acta med. acad. sci. Hung. 21 no.3:257-262 '65.

1. II. Medizinische Abteilung und Zentrallaboratorium des Bajcsy-Zsilinszky-Krankenhauses, Budapest. Submitted November 16, 1964.

HUNGARY

KOCSIS, Gyorgy; JAKO, Janos; Clinic of Dermatology and Venereal Diseases of the Medical University (Orvostudományi Egyetem Bor- és Nemibeteg Elinikája), Szeged.

"Continuous Electrophoresis."

Budapest, Kiserletes Orvostudomány, Vol 14, No 5, Oct 62, pp 535-544.

Abstract: [Authors' Hungarian summary] Modern protein research obtained many of its results by means of continuous electrophoresis. The method and the results are briefly reviewed. The authors describe their Grassmann-type apparatus, built in 1959. They also summarize their results which were obtained in experiments designed to establish their method and to reproduce data already published. [81 references, predominantly Western.]

YAKO

POLAND / Chemical Technology. Processing of Naturally Deposited Solid Fuels. H

Abs Jour: Ref Zhur-Khimiya, No 22, 1958, 75186.

Author : Yako, Takaoh, Vosatko.

Inst : Not given.

Title : Experiments in Preparing Coke From Non-Coking Coals in Hungary.

Orig Pub: Koks, smola, gaz., 1957, 2, No 6, 299-303, Diskus, 303.

Abstract: Results are reported on the preliminary experiments that were carried out in chamber furnaces (Didge's type) for producing coke from native brown coals. The experiments were varied: briquetting prior to coking, coking followed by briquetting and also repeated coking.

Card 1/2

37

AUTHOR: Jako, Ludwig SOV/68-59-5-24/25
TITLE: The Use of Coal Briquettes in Coking Charges
(Primeneniye ugol'nykh briketov v shikhte dlya
koksovaniya)
PERIODICAL: Koks i khimiya, 1959, Nr 5, pp 62-63 (USSR)
ABSTRACT: Abstracted from: Koks-Smola-Gaz, 1958, Nr 2
(Polish journal).
Abstracted by V.F. Sakhnenko.

Card 1/1

a. Jan.

- "Development and Tasks of the Innovator Movement in the Building Industry." p. 4
- "The Building of the People's Stadium Speeded with Innovations." p. 6
- "Innovators for 120,000 Dwellings." p. 7
- "Conference of Innovators in the Building Industry at Hologhassa." p. 8
- "A Criticism of the Innovator Movement in the Cement Factory in Debrecen." p. 8
- "Sheet Clippings in Electrical Engineering." p. 9
- "The Electricians Discussed their Innovation Problems." p. 10
- "New Hungarian Machines of the Building Industry Constructed Through Innovations." p. 10
- "Our Miners Following Comrade Rakosi's Teaching." p. 11
- "Results of Metallurgical Innovators in the First Quarter of the Year." p. 11
- "The Stakhanovite Innovator of the Criszologep Factory." p. 11
- "Innovation Tasks in the Mechanization of Agriculture." p. 12.
- "The Innovators Became the Representatives of our Working Peasants." p. 12
- "Istvan Machovits, a Kossuth Prize-Winning Innovator." p. 13
- "Andor Budincseviics, a Kossuth Prize-winning Innovator." p. 13
- "Stakhanovites of the Turners' Contest." p. 13
- "The Innovator Movement in Poland." p. 14
- "Soviet Building Constructions." p. 15
- "Assembly Line Production in the Building Industry." p. 15
- "A Soviet Turner as an Innovator." p. 15
- (Ujitol Lapja, Vol. 5, no. 8 Apr. 1953 Budapest.)

SO: Monthly List of East European Accessions. Vol. 2, no. 9
Library of Congress, Sept 1953, Uncl.

JAKOB, Gaon, d-r

Use of soluble antigen prepared from domestic strains of *Rickettsia prowazekii* in laboratory diagnosis of typhus. Med. arh., Sarajevo 13 no. 1:31-42 Ja-F '59.

1. Epidemioloski institut Med. fakulteta u Sarajevu, ref: prof. d-r M. Aranicki.
(TYPHUS diag.)
(ANTIGENS)

ARANICKI, Milos; JAKOB, Caon; SMJTELIV, E.

Recent epidemiology studies on endemic nephropathies in People's Republic of Bosnia and Hercegovina. Med. arh. 15 no.3:99-130 My-Je '61.

1. Epidemioloski institut Medicinskog fakulteta u Sarajevu (Sef: prof. dr Milos Aranicki) Cnetralni higijenski zavod u Sarajevu (Direktor: dr Ante Jamnicki).
(KIDNEY DISEASES epidemiol)

GOMORI, Pal; NAGY, Zoltan; JAKOB, Imre; VOJDA, Vera

On some problems related to the investigation of renal circulation.
Biol orv kozl MTA 11 no.4:383-396 '60. (EEAI 10:5)

1. Budapesti Orvostudományi Egyetem II. sz. Belklinika ja.
(KIDNEYS)

H/502/62/031/001/001/002
D409/D301

AUTHORS: Bánkóvi, Gy., Sarkadi, K., Horváth, J. and Jakob, K.
TITLE: The design and evaluation of diesel-oil desulphurization experiments by mathematical-statistical methods
SOURCE: Academia scientiarum hungaricae. Acta chimica, v. 31, no. 1-3, 1962, 23-30

TEXT: The High-Pressure Research Institute in Budapest - Pétfürdő is conducting research on hydrotreating of sulphur-rich diesel-oil cuts obtained from Soviet crude. To facilitate the tedious experiments, the mathematical-statistical method of so-called factorial experiments with partial repetition was used and is described in this article. This widely used method was slightly modified to meet the requirements of experiments aimed at determining the influence of operating conditions on the efficiency of the hydrotreating process. The test results can generally be formulated

$$z = f(u, v, x, y) + \varepsilon_{u, v, x, y}$$

Card 1/3

H/502/62/031/001/001/002
D409/D301

The design and evaluation ...

where $f(u, v, x, y)$ is the systematic influence of operating conditions (pressure, temperature, space velocity, and gas-to-product ratio), and $\varepsilon_{u, v, x, y}$ are random variables with expectation zero. Using this mathematical model and some simplifying assumptions (neglecting of higher-order interactions), it was possible to reduce hydrotreating experiments from 81, i.e. all possible combinations of the four factors in three levels, to only 36 at an estimated error (block design) of $\pm 4 - 5\%$. The hydrotreating tests proper were performed in a 200 ml laboratory-scale and a 400 l semi-production scale reactor. It was found that the desulphurization efficiency could be increased by raising the reaction temperature (to 360 - 390°C) or pressure, and reducing the space velocity. An optimum desulphurization degree was attained at a gas-to-product ratio of 500 Nm^3/m^3 . There are 2 figures and 1 table. The English-language references are: O. Kempthorne: The Design and Analysis of Experiments. (Wiley, New York) 1952; D.J. Finney: An Introduction to the Theory of Experimental Design. (The University of Chicago Press) 1960; K.A. Brownlee: Industrial Experimentation. 1947.

Card 2/3

KUCHAR, Lumir, inz., C.Sc.; BLAHOZ, Otakar, inz.; JAKOB, Miloslav, inz.

Corrosion of materials in the barite furnace. Sbornik skol ban 8
no.3:313-319 '62.

1. Odborni asistenti katedry nauky o kovech, Vysoka skola banska,
Ostrava.

JAKOŠ, Miloslav, Ing.; JAKOŠOVÁ, Anna, Ing.

Methods of corrosion measurement of the glued metal joints.
Sborník škol. báň. 8 no.3:321-327 '62.

1. Odborný asistent katedry nauky o kovech, Vysoká škola báňská,
Ostrava (for Jakob).

KUCHAR, Lumir, inz., C.Sc.; JAKOB, Miloslav, inz.

Practical use of mathematical curve analysis of aluminum alloy
metallographic diagrams. Sbor VSB Ostrava 8 No.5:545-558 '62.

1. Katedra nauky o kovech, Vysoka skola banska.

JAKOB, Miloslav, inz.; OPLEROVA, Ludmila

Hardening of leather shape-kives. Sbor VSB Ostrava 8 no.5:589-
600 '62.

1. Katedra nauky o kovech, Vysoka skola banska, Ostrava.

JAKOB, M., inž.

Formation and development of fatigue cracks. Sbor VSB
Ostrava 9 no.3:365-377 '63.

1. Katedra nauky o kovech, Vysoka skola banska, Ostrava.

TEINDL, J., prof. inž. DrSc.; KUČERA, L., inž. CSc.; JAROS, M., inž.

Causes of enamel chipping in cast-iron castings. Sbor
VSB Ostrava 9 no.3:453-466 '63.

1. Katedra nauky o kovech a tepelného zpracování, Vysoká škola báňská, Ostrava.
2. Člen korespondent Československé akademie věd (for Teindl).

JAKOB, Miloslav, inz.

Methods of determining fatigue cracks. Sbor VSB Ostrava 10 no.3:
395-402 '64.

1. Chair of Metal Science of the Higher School of Mining,
Ostrava. Submitted June 20, 1963.

JAKO, Peter, dr.

Hemangiomatosis and dyschondroplasia (Maffucci's syndrome).
Orv. hetil. 106 no.37:1759-1760 12 S'65.

1. Orszagos Testnevelési és Sportegészségügyi Intézet, Belosztály
(előíró: Lang, István, dr.).

BIRO, Andras, dr.; LOINCZ, Bela, dr.; JAKOB, Ilona, leznelked munkatars.

Our experiences with blood and fluid infusion through the
subclavian vein. Orv. hetil. 105. no.6:265-266 9 F'64

1. Baraolti Egyesitett Korhaz Sebészeti Osztaly (Roman Népköztar-
sasag, Brasov tartomány).

*

IANCU, A.; JAKOB, S.; DIVIN, M.; IANCU, A., Jr.; SURIANI, T.; VLADUTIU, V.

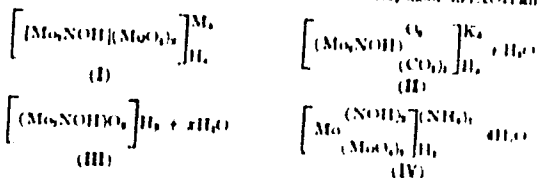
The EEG in pediatric dystrophy. Cesk. pediat. 19 no.6:528-529
Je'64.

1. Detska klinika university v Kluzi (prednosta: prof. dr. A.
Iancu); Neurochirurgicka nemocnice v Kluzi (reditel: dr. S.Jakob).

Reduction of compounds of sesivalent molybdenum by hydrazine. W. F. JAKOB AND W. KOZLOWSKI. *Russkhi Chem.* 9, 667-73 (1975 German) (1928) In the presence of NH_2 , on compounds of Mo^{VI} , NH_2 is oxidized practically completely to N_2 . NH_2 under the best conditions (high acid concn. and high temp.) reduces Mo^{VI} only to Mo^{IV} . Thus, NH_2 is a suitable reducing agent for prepn of Mo^{IV} compounds. Complete contg. Mo^{IV} and Mo^{VI} as oxidation reduction complexes were observed by partial reduction of the molybdates. The complex anions of these compounds are formed only in weakly acid soln. molybdenum blue being obtained in stronger acid soln. while in strong acid concns. the reduction of Mo^{VI} to Mo^{IV} takes place directly without the formation of the oxidation reduction complexes as intermediate products. Ammonium paramolybdate (14 g) was dissolved in 150 cc. H_2O , acidified with 3 cc. AcOH (10%), 2 g. hydrazine sulfate in 100 cc. H_2O was added and the soln. heated slowly to boiling until N_2 evolution had ceased. NH_4Cl (2 g) was added to the hot soln., the ppt. was filtered and 3 g. NH_4Cl more was added at 40°. Crystals sep'd. after 4-6 days were recrystallized from alc. giving a red brown salt, $\left[\begin{array}{c} \text{VI} \\ \text{MoO}_4 \\ \text{O} \\ \text{V} \end{array} \right] \text{NH}_4$. In an analogous way the corresponding Ba salt ($\cdot 2\text{H}_2\text{O}$) was obtained as a brown ppt. less sol. in H_2O than the NH_4 salt. P. 12

ASB 55A METALLURGICAL LITERATURE CLASSIFICATION

Compounds of hexavalent molybdenum with hydroxylamine. W. F. JAKUBASIK, B. JAKUBASIK, *Kochsche Chem. H.*, 229, 340 (German 232, 30 (1911)). Heide and Hoffmann's compounds (*Z. anorg. allgem. Chem.*, 12, 277 (1896)) prepd. by heating a molybdate with $\text{NH}_4\text{OH} \cdot \text{HCl}$ do not contain Mo of a lower valency, as some authors state, but their reducing properties and color must be ascribed to the combined NH_4OH . Analyses show that the salts have the general formula I, where all Mo atoms are hexavalent. Reduction of these salts by the iodometric method or with NH_4Ag gave no concordant results. NH_4OH in the salts of this type was detd. by decoupling of I in a 15% H_2SO_4 soln. with 4% ferric alum at the boiling temp. in a CO_2 stream. The K salt crystallizes with $14\text{H}_2\text{O}$, is a brown red microcryst. powder, probably trichloric, slightly sol. in water, sol. in dil. AcOH , sol. in strong acids with decoupling, and has a color varying with the strength of the acid; it is sol. in weak alkalis and alkali metal carbonates under decoupling. It loses $14\text{H}_2\text{O}$ at 105° , without any change in the chem. character. The NH_4 salt resembles the K salt. The Ba salt is microcryst. The Na salt (with $1 \cdot \text{H}_2\text{O}$) is prepd. from the Ba salt by interaction with Na_2SO_4 in 1% AcOH mono- or tri- clinic brownish red crystals, very sol. in water, insol. in EtOH and acetone.



By treatment of the K salt of this series with KHCO_3 , the compound II is formed, which when

treated with dil. acids liberates CO_2 and gives the free acid III. The anion is precipitated by this hydroxylaminomolybdate acid gives with alkalis deep red salt-sens, with acids intensively colored complex compds. The group contains the 10 valent nucleus Mo_2NOH . Oxidation of the hydroxylamine in this compd. with NH_4AgNO_3 is possible only in the presence of a strong base, after decomposition of the complex. Oxidation in acidic solns. yields NO as a by-product. The NH_4 salt (IV) of 1 polyhydroxylamino compd. is described. The complex of other complex higher hydroxylamine, also of that described by Cauter (C. A. 22, 1022, 24, 3723) is doubtful. Theoretically NH_4OH acts upon molybdate acid ions as follows: Polyhydroxylamino complexes are, as combinations of the oxidizer (Mo^{6+}) and the reducer (2NOH^-), an initial stage in the reduction. The true reduction process, however, takes place in the complex itself as a result of the deformation of the electronic orbits which combine the oxidizer with the reducer. Thus in the polyhydroxylamino complexes the Mo^{6+} ions are transformed into Mo^{5+} and the NOH^- ions into NOH . This deformation process is illustrated by electronic models. In the case of Heide-Hoffmann's salt, which is an oxidation product of low valent $\text{Mo-NH}_2\text{OH}$ compds., the central Mo atoms are hexavalent, but the nonsolar NOH group causes also a deformation of the electronic orbits, and hence both internal Mo^{6+} ions assume an ionic structure of a lower valency. F. Wiesner in

CA
Quadrivalent molybdenum. I. Synthesis of complex cyanides. WINTER, P.

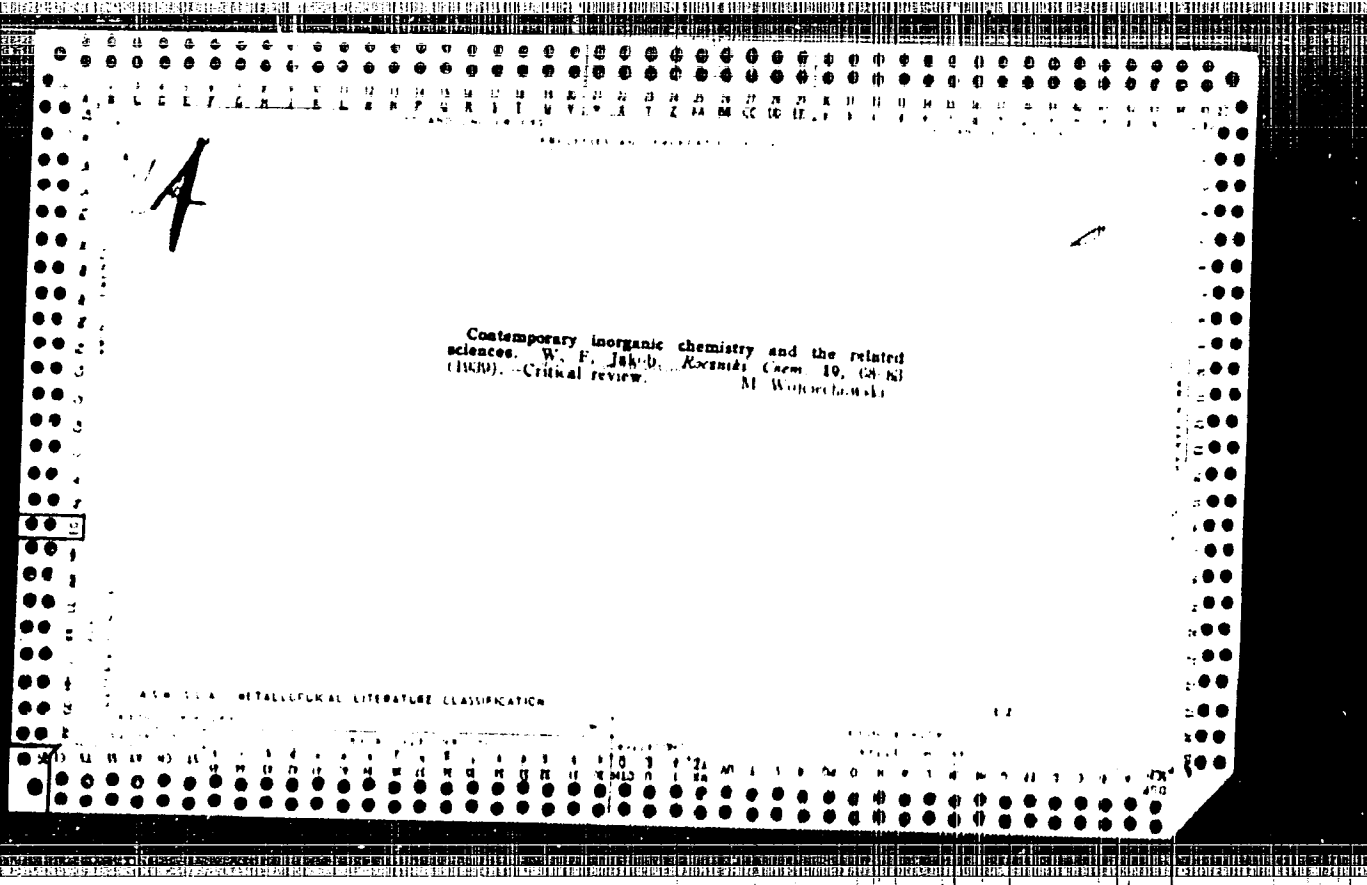
JAKOB AND EUGENIUS TUBERWICZ. *Rozwisk Chem* 11, 560-567 (1931) - The formation of $K_2Mo(OH)_4(CN)_2$ according to Bucknall and Wurdaw (1931) - The formation of Mo^{VI} to Mo^{III} and Mo^{IV} , only the latter (C. A. 22, 921) is attended by a decompos. of Mo^{VI} to Mo^{IV} and Mo^{III} in pptd. combines with KCN. To Klason's salt, $(NH_4)_2MoO_4$, neutralized with NH_3 , 2 to 4 mols. of KCN for 1 mol. of Mo is added and the mixt. is heated to 70° . Mo^{VI} is first with $BaCl_2$, the ppt. dissolved in HCl , and Mo^{VI} is detd. stannometrically. Mo^{VI} is first oxidized with $HCl + HNO_3$ to Mo^{VI} and then analyzed, as above. Prepn. of hydroxy cyanides: 100 g. NH_4 molybdate, dissolved in 100 cc. HCl , reduced with 17 g. N_2H_4 , HCl and the resulting $Mo(OH)_3$ treated with 200 g. KCN and 30 g. KOH , yields 40 g. $K_2Mo(CN)_4(OH)_2 \cdot 2H_2O$. The Na salt is prepd. in a similar manner, except that it is not pptd. with $NaOH$, but with $EtOH$. $K_2Mo(CN)_4 \cdot 2H_2O$ is prepd. by addn. of 4 mols. of KCN to a concd. soln. of the hydroxy cyanide, satn. with Cl_2 , neutralization with $AcOH$ and pptn. with $EtOH$. $Mo(OH)_3$ darkens when treated with KOH in a H_2 atm., and the filtrate contains much Mo^{VI} . The black Mo hydride is an impure hydroxide of Mo^{III} . J. WINTER

Quadrivalent molybdenum. II. Hydrolysis of complex cyanides of the type $\text{Na}_2[\text{Mo}(\text{CN})_4(\text{OH})_2]$. A hydroxide of quadrivalent molybdenum. W. F. Jakiš and C. Michalewicz. *Rozniti Chem.* 12, 570-586 (1937, 8 in English) (1932); cf. C. I. 26, 2033. The hydrolysis of red Mo hydroxycyanides proceeds in two steps and is influenced by H ions. In pure H_2O only blue products of the hydrolysis are obtained, *nr.* $\text{Na}_2[\text{Mo}(\text{CN})_4(\text{OH})_2 \cdot 2\text{H}_2\text{O}]$, blue, strongly double refracting needles, from a soln. of 10 g. of the red $\text{Na}_2[\text{Mo}(\text{CN})_4(\text{OH})_2] \cdot 12\text{H}_2\text{O}$ (I) in 150 g. H_2O with 7.5 g. HCl . $\text{K}_2[\text{Mo}(\text{CN})_4(\text{OH})_2]$ results from the neutralization of the red $\text{K}_2[\text{Mo}(\text{CN})_4(\text{OH})_2] \cdot 6\text{H}_2\text{O}$ (II) with CO_2 , AcOH or NH_4OAc . *Cd salt*, $[\text{Cd}(\text{H}_2\text{O})_6][\text{Mo}(\text{CN})_4(\text{OH})_2]$, blue-purple, from neutralization of I with a 1% soln. of AcOH and addn. of CdCl_2 . *Ammonio Cd salt*, $[\text{Cd}(\text{NH}_3)_6][\text{Mo}(\text{CN})_4(\text{OH})_2]$, purple crystals, insol. in H_2O , sol. with blue color in concd. NH_3 , from the interaction of the red alkali salts and an NH_3 soln. of CdCl_2 in presence of NH_4Cl . It is decomposed by hot Na_2CO_3 soln. with evolution of NH_3 and formation of CdCO_3 . *Mn salt*, $[\text{Mn}(\text{H}_2\text{O})_6][\text{Mo}(\text{CN})_4(\text{OH})_2]$, blue-purple crystals, from neutralization of I and addn. of MnCl_2 . *Ammonio Mn salt*, purple ppt., $[\text{Mn}(\text{NH}_3)_6][\text{Mo}(\text{CN})_4(\text{OH})_2] \cdot \text{H}_2\text{O}$, from the addn. of MnCl_2 and NH_3 to the nearly neutralized soln. of II. In the presence of larger amts. of NH_3 another salt, richer in NH_3 , is formed: $[\text{Mn}(\text{NH}_3)_6][\text{Mo}(\text{CN})_4(\text{OH})_2] \cdot \text{H}_2\text{O}$. The solns. of I and II become green on addn. of even the weakest acids, especially if heated, whereby gels are formed contg. less CN than the original salts. II does not become blue on keeping over solid KOH or CaCl_2 , but does so in the presence of moisture or acidic vapors. I is more readily decompd. than II. The bimetallic salts are more effectively hydrolyzed only in the presence of H ions: $[\text{Mo}(\text{CN})_4(\text{OH})_2]^{2-} + 2\text{H}^+ \rightarrow \text{Mo}(\text{CN})_4(\text{OH})_2(\text{H}) + 2\text{HCN}$. III, a dark-green gel, shows no acidic properties. It is peptized by the action of bases and, being unstable, it is converted irreversibly into $\text{Mo}(\text{OH})_4$. The latter

can be prepd. also by pptn. with alkali from the product of reaction of I or II with concd. HCl. The gel is red-brown in transmitted, green-brown in reflected light, and is oxidized by air in the presence of alkalis. Purified with NH_4Cl , H_2O and Et_2O it shows the compn. $\text{MoO}_3 \cdot 11\text{H}_2\text{O}$; it is readily sol. in concd. acids, its solns. are red to brown-purple. Its acid solns. have a weaker reducing power than similar solns. of Mo^{VI} or Mo^{III} compds. The potential of a Pt electrode in acid solns. is $E = 0.27$ v. A jump corresponding to the intermediate transition of Mo^{VI} into Mo^{IV} during the KMnO_4 titration of Mo^{VI} solns. could not be observed, and hence it appears that the compd. is oxidized directly to Mo^{VI} .

J. Wertelak

The influence of complex formation on the attainment of equilibrium in some oxidation-reduction systems. WIKTOR F. JAKOŃ AND MARJAN R. RUSNAK, *Chem. Listy* 26, 101 (1931) (in Polish). *Collection Czechoslov. Chem. Commun.* 6, 303 (1931) (in English). Solns. of $[H_2MoO_4]_2$ and H_2SO_4 (I) were pipetted into weighed quantities of $NH_4MoO_4 \cdot 2H_2O$ (II) and equal potentials were noted in a stream of CO_2 . The pH was maintained const. (≈ 0.2) with a large excess of acetate buffer. The stream of CO_2 showed no change in acidity of the soln., buffer mixts. of the same acidity had no noticeable effect on the oxidation-reduction potentials. The curves obtained were characteristic for all oxidation-reduction systems obeying the Nernst law, except at low acidities, where more complicated phenomena are taking place, and the curve deviates from a logarithmic form. The pure complex II imparted a large potential to the indifferent electrode, but with increasing contents of Mo the potential equally increased in the direction of the noble potentials. The anions of the complex II function as an active reducing agent, the Mo and H ions play the role of oxidizing agents toward them. A considerable sensitivity of the electrode toward small additions of Mo to weakly acidified solns. of II may indicate a slight hydrolysis of the oxidation-reduction complex and liberation of Mo acid ions. To prep. II dissolve 11 g. NH_4 molybdate in 100 cc. H_2O contg. 3 cc. 50% $AcOH$, add to 2 g. hydrazine sulfate in 100 cc. H_2O , heat until the evolution of N ceases, add to the hot soln. 2 g. NH_4Cl , filter, cool to 40° , treat with 2 g. NH_4Cl , after 48 hrs. decant the dark ruby crystals from the slime, wash with 50, 50 and 90% $EtOH$ and with ether, and dry in air. I was prepd. by crystg. the com. form from weak NH_4 solns. of $NH_4H_2MoO_4$ and $MoO_3 \cdot 2H_2O$ was prepd. from partially reduced Mo solns. of molybdenum green compound is crystals as dark blue crystals, the crystals for a few days, which through hydrolysis changes through green to a light brown.



Quadrivalent molybdenum. III. Dioxochloromolybdous acid. Stability of acid solutions of quadrivalent molybdenum. W. P. Jakób and L. Czirny-Schulowski. *Rozprawy Chem.* 10, 116-120 (1930); cf. C. A. 27, 3228. $K_2(MoO_2)_2(OH)_2$ heated with dil. HCl yields $Mo(CN)_2(OH)_2$, which is boiled under reflux (3 hrs.) with concd. HCl. The soln. is concd. in vacuo to a syrup, which is extrd. with Et_2O . This dissolves $H_2MoO_4 \cdot 2H_2O$, leaving H_2MoCl_4 in the aq. layer, from which a violet oil seps., yielding solid $MoCl_4(OH) \cdot 3H_2O$ (1) when dried. Solns. of I are violet, yield a brown ppt. with aq. NH_3 , and do not change color with CNS^- or MoO_4^{2-} . IV. Decomposition of octacyanomolybdous acid. Dicyanic acids. *Ibid.* 151-5. — $K_2(Mo(CN)_2)_4$ boiled with 3% H_2SO_4 yields HCN and $Mo(CN)_2(OH)_2 \cdot 3H_2O$, oxidized by H_2O_2 to $H_2Mo(CN)_2 \cdot 3MoO_4 \cdot 3H_2O$. H. C. P. A.

ADP 31.6 METALLURGICAL LITERATURE CLASSIFICATION

COMMON ELEMENTS										PROPERTIES AND PROPERTIES INDEX									
<p>2</p> <p>/ Modern atomic weights. Wiktor Jakob (illegible) Politechn. School, Gdansk, Poland). <i>Prilozhenie</i> 4. 20-24 (1940).—An address. A. Bogoyavl</p>										<p>2</p>									
<p>ASS-ILA METALLURGICAL LITERATURE CLASSIFICATION</p>										<p>W-2 272-24121</p>									
<p>10000 #1</p>										<p>10000 #1</p>									
<p>10000 #1</p>										<p>10000 #1</p>									

Coordination number ten in free complex ions. W. J. I. J. and Z. J. J. (U.S. Pat. 2,644,444, 1954).
 "Chem. Abstr. 49:1162 (German summary) --
 $[W(CN)_4] \cdot 4H_2O$ and $M[Mo(CN)_4] \cdot 4H_2O$, in which M
 is Cd or Mn, and R is H_2O , NH_3 , or NH_4 , were prepared.
 The NH_3 and NH_4 compds. are particularly stable. They
 sep. as insol. red crystals from aq. soln. Conductometric
 measurements on aq. complexes of the type $[M(CN)_4]^{4-}$
 indicate that the coordinated groups H_2O , NH_3 , and NH_4
 are a part of the free neg. ions $[Mo(CN)_4]^{4-}$ and
 $[W(CN)_4]^{4-}$. The aq. solns. of these are stable in the
 dark, and are hydrolyzed in light to $[Mo(CN)_4(OH)]^{3-}$
 and $[W(CN)_4(OH)]^{3-}$.
 Michael Perl

Distr: 4E2c

Photochemical reactions of octacyanides of molybdenum (IV). Zbigniew Jakób and Wiktor Jakób (Univ. Kraków, Poland). *Zeszyty Nauk. Inst. Chem. PAN* 1971, Prayrod., Mat., Fiz., Chem. No. 2, 40-60 (1980) (English summary).— $K_2Mo(CN)_8$ (I) was prepd. by the modified method of W. Jakób and Turkiewicz (C.A. 26, 2934a). The procedure is: Reduce MoO_3 with excess hydrazine sulfate (II) in hot concd. HCl (1.5 ml./g. MoO_3), filter the red-brown soln., dil. with large amt. of H_2O , ppt. $MoO(OH)_3$ with a small excess NH_3 , wash, filter, add 2.5 molen KCN per 1 mole Mo, heat, and add 0.25 mole KOH, evap. H_2O in *vacuo*; when blue crystals appear add further small portions of KOH, cool, and filter the red-brown $K_2Mo(CN)_8 \cdot (OH)_2$ (III); expose the green filtrate to light, filter, and combine the 2 portions of III. Add 1 mole III to 1 l. 0.2N

KCN, sat. with CO_2 with vigorous shaking, when yellow or brown color appears, neutralize with concd. AcOH passing a stream of air through the soln., evap. in *vacuo*, filter, and wash the resulting I twice with 50% and twice with 90% EtOH. Yellow I (5 g. $1.2H_2O$ in 1.6 l. H_2O), exposed to daylight at 14-17°, became orange, red, and violet. In all cases only III was isolated, contrary to Collinberg (C.A. 18, 3323). After 45 min. the red color intensity reached a max., and upon interruption of exposure yellow I was regenerated. Violet solns. afforded either $III \cdot 6H_2O$ upon KOH addn., or violet $Cd(NH_4)_2Mo(CN)_8(OH)_2$ upon Cd^{++} , NH_4Cl , and NH_3 addn (C.A. 27, 5019). No photolysis was detected at 40° and above. From nonirradiated I, cryst., sparingly sol., yellow $Cd_2Mo(CN)_8 \cdot 8H_2O$, yellow $Mn_2Mo(CN)_8 \cdot 8H_2O$, and dark-yellow $TlMo(CN)_8$ were obtained. To 1.5 l. aq. soln., contg. 5 g. $1.2H_2O$ and 80 ml. 2N NH_3 , irradiated to brown-red, 60 ml. 0.5N $CdNO_3$ was added; cryst. red $Cd_2Mo(CN)_8(NH_4)_2 \cdot 4H_2O$ was obtained.

CC
1/2

POLAND/Inorganic Chemistry - Complex Compounds.

C.

Abs Jour : Ref Zhur - Khimiya, No 11, 1958, 35675

Author : Jakob Wiktor, Ogorzalek Maria

Inst : -

Title : The Nature of Peroxidation Bridges in Binuclear Cobalt-Ammine.

Orig Pub : Roczn. Chem., 1956, 30, No 4, 1055-1066

Abstract : The decomposition process of I in an alkali medium has been investigated in order to explain the structure of the complex $[\text{Co}_2\text{O}_2(\text{NH}_3)_{10}]^{5+}$ (I). The reaction between the solid phase $[\text{Co}_2\text{O}_2(\text{NH}_3)_{10}](\text{NO}_3)_4 \cdot \text{H}_2\text{O}$ and a HNO_3 solution has also been studied. This reaction proceeds according to the composite equation: $6 [\text{Co}_2\text{O}_2(\text{NH}_3)_{10}]^{4+} + 10\text{H}_2\text{O} = 2\text{I} + 8 [\text{Co}(\text{NH}_3)_5 \text{H}_2\text{O}]^{3+} + 7\text{H}_2\text{O} + 3/2 \text{O}_2$.

Card 1/2

$[\text{Co}(\text{NH}_3)_5 \text{H}_2\text{O}]^{3+}$.

Card 2/2

JAKOB, Wiktor; SANDTUS-KOSINSKA, Alina; STASICKA, Zofia

On investigations of the photochemical reactions of octacyano-
molybdates (IV) and octacyano-tungstates (IV). Roczniki chemii
36 no.1:165-167 '62.

1. Department of Inorganic Chemistry, Jagellonian University,
Krakow.

JAKOB, Wiktor; JAKOB, Zbigniew [deceased]

Investigations of the photochemical reactions of octacyanomolybdates (IV) and octacyanotungstates (IV). Pts. 1-2. Pocz chemii 36 no.4: 593-609 '62.

1. Department of Inorganic Chemistry, Jagellonian University, Krakow.

MAKOB, Wiktor, prof. dr

Dr. Jan Zygmunt Robel; obituary. Wiad chem 17 no.6:321-324
Je '63.

1. Kierownik Zakladu Chemii Nieorganicznej, Uniwersytet
Jagiellonski, Krakow.

INORGANIC Chem

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C. 62.

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OP JAKOB, Z. L.

Theory of acidimetric analysis. Z. L. Jakob (Warsaw, Poland). *Bull. intern. acad. polon. sci., Cl. III sci. math. et nat.*, Ser. A, 1950, 70 800 (English). Roller's equations (C. A. 44, 8273) for errors in acidimetry are modified to the form $E = 100\sqrt{K/C_0}(10^{\Delta pH} - 10^{-\Delta pH})$ and $\Delta pH = P_i - P_s + \Sigma a$, in which E = % pH uncertainty error, K = eq. const. in the titrated soln., C_0 = final concn. of the product of titration, P_i = acidity indicated by the indicator, P_s = stoichiometric acidity after titration, Σa = sum of empirical corrections for salt and colloidal effects on the indicator and the uncertainty in detecting the color change. 1 P. 8

1952

CA 4450, 50

Errors in acidimetry and alkalimetry. Zbigniew L.
Jakob (Higher Polytech. School, Gliwice, Poland). *Anal.*
Chem. 4, 305-16(1950). An address. A. S.

JAFI 010, 2.

JAFI 010, 2. Field of antenna for directing ultra short wave communications. p. 108.

Vol. 9, No. 10/11, 1958

ELFETECNICAR.

TECHNOLOGY

Zagreb, Yugoslavia

See: East European Acquisitions, Vol. 5, No. 5, May 1958

JAKOBCZYK, F. (Lublin)

On certain properties of the functions $h_q(n)$ and $L_q(n)$ and their application to the study of periodicity of the series $\{q^n\}_{\text{mod } m^k}$ ($n = 1, 2, 3, \dots$). Annales pol math 9 no.1:1-24 '60.

(Zbl 10:9/10)

(Numbers, Theory of) (Functions) (Series)

S/274/63/000/002/007/019
A055/A126

AUTHORS: Martyniuk-Lewko, Sergiusz, Jakóbczyk, Mieczysław

TITLE: Time-sweep generator

PERIODICAL: Referativnyy zhurnal, Radiotekhnika i Elektrosvyaz', no. 2, 1963,
63, 2A385 P (Polish pat., cl. 21 e, 28/02, no. 44342, April 10,
1961)

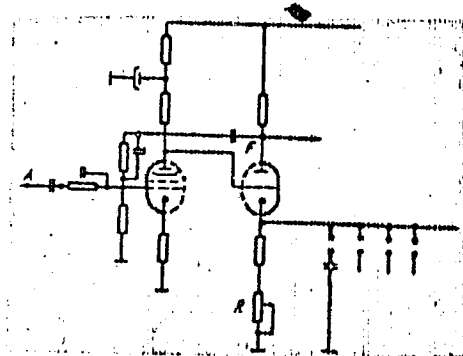
TEXT: The object of the patent is a horizontal sweep generator circuit for oscillographs (see Fig.), consisting of a pentode preamplifier and an output stage with anode-cathode load, with strong positive feedback. The cathode load of the output stage is shunted by a capacitor, whose value varies depending on the position of the range-switch; a continuous frequency-control is obtained by means of the variable resistance R in the output stage cathode. The synchronization signal is applied to the terminal A.

Card 1/2

Time-sweep generator

S/274/63/000/002/007/019

Figure



I.Z.

[Abstracter's note: Complete translation]

Card 2/2

JAKUBOWSKI, B.

Source: given names

Country: Poland

Academic Degrees: /not given/

Affiliation: /not given/

Source: Warsaw, Medycyna Weterynaryjna, Vol XVII, No 6, June 1961, p 338.

Data: "Increased Control of Trichinellosis."

JAKOBIEC, M.

Diagnostic difficulties and therapeutic results of streptomycin
in adrenal cortex insufficiency. Polski tygod. lek. 7 no.1-2:34-
38 7 Jan 1952, (CLML 22:2)

1. Of the First Clinic of Internal Diseases (Head--Prof. Leon
Tochowicz, M. D.) of Krakow Medical Academy.

JAKOBIŃC, M.

A case of typhoid fever bacilli carrier treated by chloromycetin.
Polski tygod. lek. 7 no.3-4:88-89 21 Jan 1952, (GIML 22:2)

1. Of the First Clinic of Internal Diseases (Head--Prof. L. T.
Tochowicz, M. D.) of Krakow Medical Academy.

JAKOBIEC, M.

Psychoneurosis as a cause of somatic emanation. Polski tygod. lek.
8 no.10:382-385 9 Mar 1953. (CJML 24:5)

1. Of the First Internal Clinic (Head--Prof. Leon Tochowicz, M.D.) of
Krakow Medical Academy.

JAKOBIEC, Mieczyslaw; KRAUSS-ZAKI, Janina

Treatment of parenchymatous jaundice with BAL. Polski tygod. lek.
9 no.26:812-814 26 June 54.

1. Z I Kliniki Chorob Wewnętrznych A.M. w Krakowie, kierownik:
prof. dr Leon Tochowicz.

(HEPATITIS, INFECTIOUS, therapy,
dimercaprol)

(DIMERCAPROL, therapeutic use,
hepatitis, infect.)

JAKOBIEC, Mieczyslaw

Inflammatory diseases of the kidneys and their treatment with systemic antibodies. Polskie arch. med. wewn. 26 no.3:347-358 1956.

1. Z I Kliniki Chorob Wewnętrznych A.M. w Krakowie, Kierownik: prof. dr. med. L. Tochowicz, Krakow, I Klinika Chorob Wewnętrznych A.M. Kopernika 17.

(GLOMERULONEPHRITIS, therapy,
urinary antibodies (Pol))

(ANTIGENS AND ANTIBODIES,

urinary antibodies, ther. of glomerulonephritis (Pol))

(URINE,

antibodies, ther. of glomerulonephritis (Pol))

"APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619420007-2

APPROVED FOR RELEASE: 08/10/2001

CIA-RDP86-00513R000619420007-2"

/Preparation of 2-aminothiazole. Bogdanov, P. Ivan K.
Todorov, L. Koldjev, and Ivan G. Petrovski. (Mol. Cryst. Liq. Cryst., 1964, 109, 271-274).
Reaction of 2,4-dinitrophenol (I) with 2,5-diaminobenzaldehyde (II) and ribonol (III). Litch and Beckman, U.S.
2,250,792 (C.A. 35, 5270) gave a very brown mass was obtained
in yield of I, presumably because, in the absence of
water, II did not depolymerize to react with III. The syn-
thesis was modified as follows: 24.8 ml. Br was added dur-
ing 3 hrs. with stirring to 30 g. paraaldehyde and 120 ml.
water with the temp. kept at 33-35°, the colorless mixt.
treated with 30 g. III, and stirring continued 1 hrs. at 75-
89°; neutralization with 50% NaOH (about 140 ml.), to
litmus at 35°, extn. with five 20-ml. portions of Et₂O, drying
with K₂CO₃, and distn. at 15 mm. gave 3g. 40 g. (60%) pure
I, bp 90°.
Janine R. Spencer

At 311

✓ Utilization of sulfate turpentine for the preparation of medicinal products. B. Bobrański, T. Jakóbczyk, and J. Pomorski (Zakład Chem. Farm. A.M., Wrocław). *Acta Polon. Pharm.* 12, 91-6(1955).--By fractional distn. of sulfate turpentine, a waste product of the cellulose industry, the sample yielded approx. 40% pinene; b. 154-60°, of sufficient purity to be used for camphor and terpene hydrate synthesis.

L. J. Piotrowski

(2)

A novel synthesis of bisethobromide of methylbis(di-
methylaminoethyl)amine. R. Bobrowski, T. Jakubiec,
and D. Pielicz (Inst. Pharm. Chem., Wrocław, Poland).
Acta Polon. Pharm. 12, 105-11 (1955) (Engl. summary);
cf. C.A. 46, 896i. —(HOCH₂CH₂)₂NH (53.5 g.) mixed with
450 ml. HBr (d. 1.473) is distd. through a 30 cm. Widmer
column until 120 ml. distillate is collected. The mixt. is
refluxed 1 hr., 155 ml. distd. off, again refluxed 3-4 hrs., 135
ml. distd. off, and the residue cooled and crystd. by adding
75 ml. AcOMe to give 102-10 g. crude NH(CH₂CH₂Br)₂.
HBr (I). 1 (30 g.), 10 g. 92% HCO₂H, and 20 ml. 35%
HCHO heated 1.5-2 hrs. yields on evapn. in vacuo 31 g.
crude MeN(CH₂CH₂Br)₂ (II), m. 147° (from AcOH-Pt₂O).
II (3.28 g.), 2.5 g. EtMe₃N, and 35 ml. abs. EtOH heated 3
hrs. yield after evapn. and addn. of 80-100 ml. abs. Et₂O
3.5 g. of MeN(CH₂CH₂NMe₂EtBr)₂.
R. Dowlingko.

dent 3

1700

PM 1955

BOBRANSKI, B.; JAKOBIEC, T.; PRELICZ, D.

New neurotropic barbituric acid derivatives. Acta Poloniae
pharm. 12 no.4:237-240 1955.

1. Z Instytutu Immunologii i Terapii Doswiadczałnej PAN im.
L.Hirszfelda. Z Zakładu Chemii Farmaceutycznej oraz II Kliniki
Chorob Wewnętrznych we Wrocławiu.

(BARBITURATES.

pharmacol. of several barbituric acid deriv.)

Country : POLAND
 Category : Organic Chemistry. Synthetic Organic Chemistry G
 Ref. Jour : Ref Zhur - Khim., No 5, 1959, No. 15432
 Author : Bobranski, B.; Jakobiec, T.; Prolicz, D.
 Institut. : -
 Title : On the Action of Iodine on 5,5-Diallylbarbituric Acid
 Orig. Pub. : Roczn. chem., 1956, 30, No 2, 483-492
 Abstract : In continuation of the work begun earlier (see report I, Ref Zhur-Khim, 1957, 19216), the structure of the product which is formed under the action of I_2 in the absence of HI on 5,5-diallylbarbituric acid (I), both in an acid and in an alkaline medium, was examined. The product obtained differed in composition from the earlier-prepared I under the action of I_2 on I in a weak alkaline medium (Bougault, J., Guillou, J., C. r. Acad. sci., 1931, 193, 463).
 Card: 1/9

G - 60

Country :
 Category :
 No. Jour : Ref Zhur - Khim., No 5, 1959, No. 15432
 Author :
 Institut. :
 Title :
 Orig. Pub. :
 Abstract : of HIO on 5-allyl-5-(β -oxy- γ -iodopropyl)-bar-
 cont'd. bituric acid (III). During the reduction of II
 with Zn powder, I is again recovered. The
 structure of II is also confirmed by the fact
 the HIO convert 5-allyl-5-(β -oxypropyl)-bar-
 bituric acid (IV) into (V), and 5-acetonyl-5-

$$\begin{array}{c}
 \text{O} = \begin{array}{l} \text{NH} - \text{CO} \\ \text{NH} - \text{CO} \end{array} \begin{array}{c} \diagup \\ \diagdown \end{array} \begin{array}{c} \text{R} \\ \text{O} \\ \text{CH}_2\text{I} \end{array}
 \end{array}
 \quad \begin{array}{l} \text{II R} = \text{CH}_2\text{I} \\ \text{V R} = \text{CH}_3 \end{array}$$

Cord: 3/9

G - 61

Category : G
 Abs. Jour : Ref Zhur - Khim., No 5, 1959, No. 15432
 Author :
 Institut. :
 Title :

Orig. Pub. :

Abstract : is dissolved in a small quantity of alcohol;
 cont'd. an aqueous solution of $\text{Na}_2\text{S}_2\text{O}_3$ is added, and
 12 g. of II is obtained, m.p. 215-218° (decom-
 position; from alcohol). Analogous results are
 obtained by conducting the reaction at diffe-
 rent values of pH > 7. 3.5 g. of III, 100 ml.
 of water, 20 ml. of 10% H_2SO_4 and 0.72 g. of
 KIO_3 are heated to 80°, 1.1 g. of KI in 20 ml.
 of water are added, and 3.5 g. of II is ob-
 tained, m.p. 214-216° (from aqueous alcohol).

Card: 5/9

G - 62

Country : G
Category :
Abs. Jour : Ref Zhur - Khim., No 5, 1959, No. 15432
Author :
Institut. :
Title :
Orig. Pub. :
Abstract : 1 g. of II, 100 ml. of water and 1 g. of Zn
cont'd. powder are boiled for two hours, and 0.3 g. of
I is separated out from the filtrate. 1.8 g.
of KI and 0.72 g. of KIO_3 in 30 ml. of water
are added to 2.3 g. of IV and 1 g. of KI in
5 ml. of hot water and 2 ml. of 16% H_2SO_4 at
 80° , washed with $Na_2S_2O_3$ solution after about
12 hours, and 2.5 g. of V is obtained, m.p.
210.5-211 $^\circ$ (decomposition; from alcohol).
2.2 g. of IV, 0.75 g. of KIO_3 , 2 ml. of 16%

Card: 6/9

Country :
Category :

Abs. Jour : Ref Zhur - Khim., No 5, 1959,

No. 15432

Author :
Instit. :
Title :

Orig. Pub. :

Abstract
cont'd.

: H_2SO_4 and 10 ml. of water are heated to 80° ,
1.1 g. of KI in 20 ml. of water are added, 2.2
g. of V is obtained, m.p. $211-212^\circ$ (decomposi-
tion; from water). 11 g. of VI, 3.6 g. of KIO_3 ,
200 ml. of water and 50 ml. of 10% H_2SO_4 are
heated to 80° , 5.5 g. of KI in 70 ml. of water
are added, and after 24 hours 12 g. of VII are
obtained, m.p. $211-212^\circ$ (decomposition; from
water); 24-dinitrophenylhydrazones, m.p. $230-$
 232° . 6 g. of VII in 250 ml. of 10% H_2SO_4 are

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7/9

G - 63

G - 64

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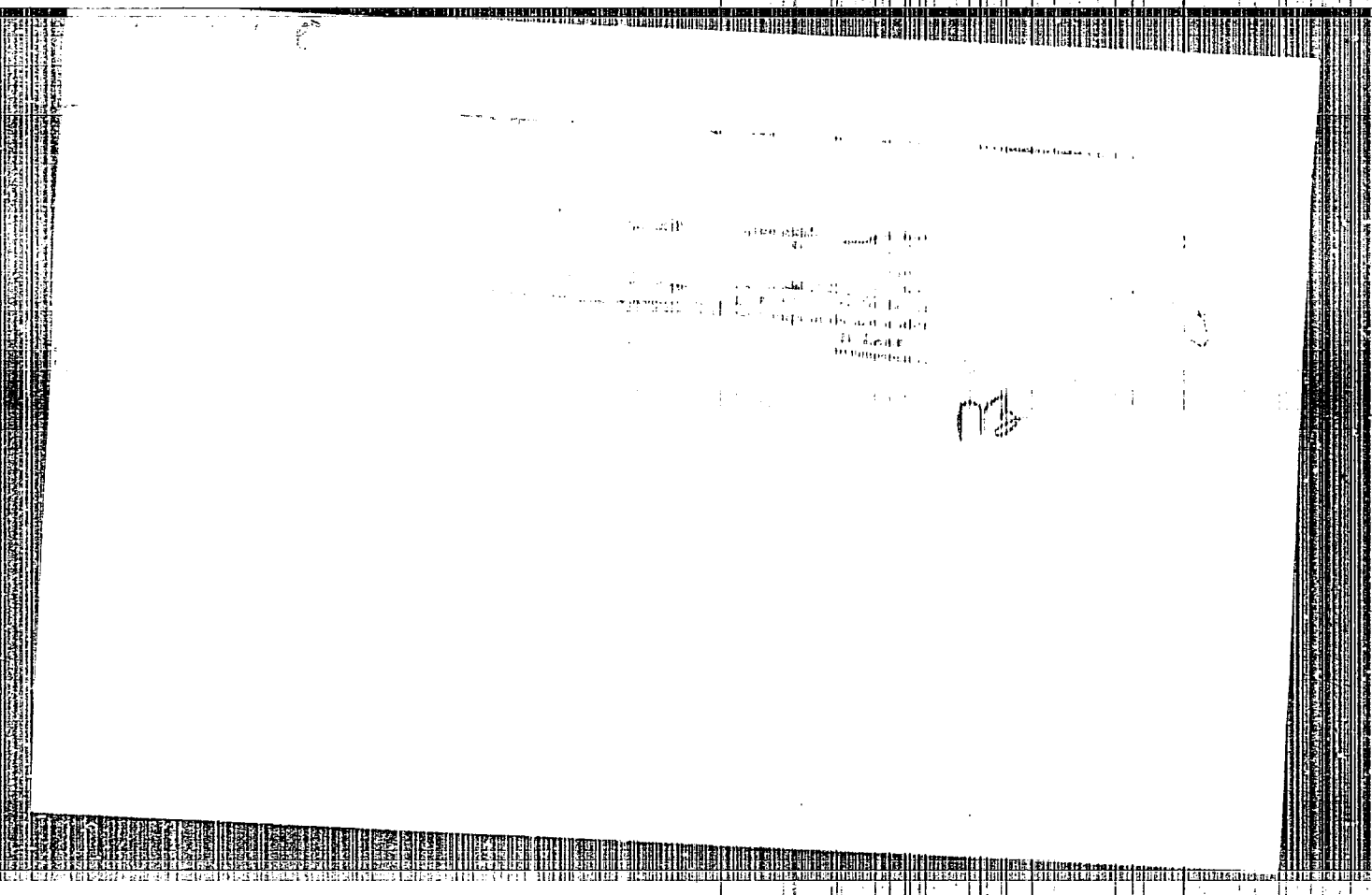
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JAKOBIEC, T.

POLAND/Organic Chemistry. Synthetic Organic Chemistry.

E-2

Abs Jour: Ref Zhur-Khimiya, No 6, 1957, 19217

Author : Bobranski B., Jakobieć T., Prolicz D.

Inst : *Res. Lab. Pharmaceutical Chem., Acad. Med. Warsaw - Inst. Immunology & Experimental*

Title : Action of Iodine on 5-isopropyl-5-allylbarbituric acid. *Therapy, 8 Polish Acad. Sci., Warsaw*

Orig Pub: Roczn. Chem., 1956, 30, No 1, 165-174.

Abstract: In quest of nontoxic preparations, having an effect on the nervous system, the reaction of iodine with 5-isopropyl-5-allylbarbituric acid (I) was studied. As a result 5-isopropyl 5-(β -hydroxy- γ -iodopropyl)-barbituric acid (II) is formed. Structure II is confirmed: 1) by oxidation with $K_2Cr_2O_7$ in an acid medium with the formation of 5-isopropyl-5-(γ -iodoacetyl)-barbituric acid (III); 2) Regeneration of I by boiling II with water and Zn-dust. III when boiled with water and Zn-dust is transformed into 5-isopropyl-5-acetylbarbituric acid

Card : 1/3

POLAND/Organic Chemistry. Synthetic Organic Chemistry.

APPROVED FOR RELEASE: 08/10/2001 CIA-RDP86-00513R000619420007-2

Abs Jour: Ref Zhur-Khimiya, No 6, 1957, 19217

H_2SO_4 is acidified with $K_2Cr_2O_7$ in 40 cc water (heating on a water bath 15 min.), and obtained are 4.5 g. III, m.p. 200-201° (dec.; from alc.); 2,4-dinitrophenylhydrazones, does not melt up to 300°. 5 g. I is dissolved in 25 g. conc. H_2SO_4 , after 15 min. it is poured into water, and obtained are 5 g. V, m.p. 188-190° (from alc.); benzoyl derivative, m.p. 173-175° (from ethylacetate); acetyl derivative, m.p. 144-145° (from benzene). 2 g. III is boiled 2.5 hours with 2 g. Zn-dust and 100 cc water and obtained are 0.5 g. IV, m.p. 259-261°; 2,4-dinitrophenylhydrazones, decomp. p. 260°. 0.5 g. V is oxidized in the same way as II, and is obtained 0.3 g. IV.

Card : 3/3

W. A. Kobielski, I.

POLAND/Organic Chemistry. Synthetic Organic Chemistry.

E-2

Abs Jour: Ref Zhur-Khimiya, No 6, 1957, 19216.

Author : Bobranski B., ~~Jakobina T.~~ Prolicz D.

Inst :

Title : Action of Iodine on 5,5-diallylbarbituric Acid. I.

Orig Pub: Roczn. Chem., 1956, 30, No 1, 175-184.

Abstract: At the action of iodine on 5,5-diallylbarbituric acid (I) in an acidic medium even with a surplus of iodine 5 allyl-5-(2-hydroxy-γ-iodopropyl)-barbituric acid only (II) is obtained. Only in the presence of a surplus of KIO_3 is the compound $C_{10}H_{12}O_4N_2J_2$ (III) obtained. The structure of II is determined: 1) by oxidation with $K_2Cr_2O_7$ in acidulous media with the formation of 5-allyl-5-(10-deacetonyl)-barbituric acid (IV); 2) the reduction of II by boiling with water and Zn-dust with the formation of I; in analogical conditions IV yields 5-allyl-

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Country : POLAND

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Category: Pharmacology. Toxicology. Ganglionic Blocking Agents.

Abs Jour: RZhBiol., No 6, 1959, No 27769

Author : Bobranski, Boguslaw; Jakobiec, Tadeusz; Prelicz, Danuta

Inst : -

Title : On New Chemical Compounds which Block the Activity of Autonomous Nerve Ganglia.

Orig Pub: Dissert. pharmoc. PAN, 1956, 8, No 4, 249-255

Abstract: Bis-quaternary nitrogenous bases of the type of pendionide are obtained by means of reacting of methyl-bis (beta-bromoethyl)-amine with tertiary amines. Compounds which contain diethylmethylamine, N-methylpiperidine, N-methylmorpholine and

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Methods of corrosion measurement of the glued metal joints.
Sbornik skol ban 8 no.3:321-327 '62.

1. Odborný asistent katedry nauky o kovech, Vysoká škola báňská,
Ostrava (for Jakob).

1. 60259-65 EMP(w)/ENA(d)/T/EMP(t)/EMP(z)/EMP(b) MJW/JD
 CZ/0032/014/014/012/0918/0928
 ABSTRACT NR: APSC19909

AUTHOR: Branka, T. (Engineer, Poldyna, V. (Engineer, Candidate of sciences))
 Jakubova, A. (Engineer)

TITLE: Heat resistance of Czech boiler steels 15110, 15111, 15125, and 15225

ABSTRACT: Strojitelnost, v. 14, no. 12, 1964, 918-928

Heat resistance of alloy steel, metal creep, metal test, ferritic steel, pearlitic steel, 15110 steel, 15111 steel, 15125 steel, 15225 steel

The article describes the results of long-term creep tests of low-alloy ferritic and pearlitic steels 15110, 15111, 15125, and 15225. The tests were conducted at temperatures of 400°C and 450°C. The results show that the steels exhibit good heat resistance and creep strength. The article also discusses the influence of microstructure and alloying elements on the creep properties of the steels.

Key words: creep, metal test, metal creep, metal test, ferritic steel, pearlitic steel, 15110 steel, 15111 steel, 15125 steel, 15225 steel

ASSOCIATION: Vyzkumny ustav metalurgicky, VZUG, Ostrava Metallurgical Research Institute, VZUG/
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